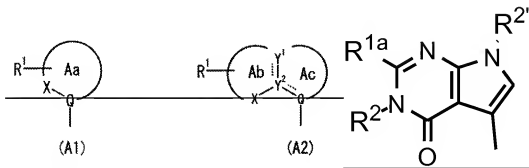


AMENDMENTS TO THE CLAIMS

1. (Currently amended) A compound represented by the formula:



wherein, A is a group represented by the formula (A1) or (A2):



wherein, ring Aa is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Q and X, and may be further substituted with one or more substituents in addition to R¹;

ring Ab is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Y¹, Y² and X, and may be further substituted with one or more substituents in addition to R¹;

ring Ac is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Y¹, Y² and Q, and may be substituted with one or more substituents;

R¹ is an optionally substituted hydrocarbyl, a substituted amino, an optionally substituted cyclic amino, a substituted hydroxy, a substituted sulfanyl, an optionally substituted sulfinyl, or an optionally substituted sulfonyl;

X is carbonyl, -O-, -S-, -SO-, or -SO₂-;

Y¹, Y² and Q are independently optionally substituted carbon or nitrogen;

--- is a single or double bond;

wherein R^{1a} is

(1) an amino which is mono- or di-substituted with

(i) a C₁₋₈ alkyl which may be substituted with a hydroxyl substituted with a C₁₋₈ alkyl, a C₃₋₇ cycloalkyl, a phenyl, a 4-methylphenyl, a hydroxyl substituted with a phenyl,

a 2-chlorophenyl, a heterocyclic group, a 4-chlorophenyl, a 4-(benzyloxy)phenyl,
a 3-methoxyphenyl, a 3-chlorophenyl, a 2'-cyanobiphenyl, a naphthyl, a 2,5-dimethoxyphenyl,
a 3-fluoro-5-(trifluoromethyl)phenyl, an acyl, or an esterified or amidated carboxyl,

- (ii) a C₂₋₈ alkenyl,
(iii) a C₁₋₁₀ acyl, or
(iv) a C₃₋₇ cycloalkyl, or

(2) a cyclic amino;

R² is a hydrogen, a C₁₋₈ alkyl which may be substituted by a cyano or a phenyl;

R^{2'} is

(1) a hydrogen,

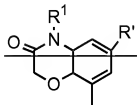
(2) an acetyl, or

(3) a C₁₋₈ alkyl which may be substituted with a phenyl, a 4-methoxyphenyl or an acetyl;

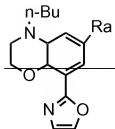
W is a bond, an optionally-substituted methylene, an optionally-substituted ethylene, an
optionally-substituted imino, O, S, SO, or SO₂;

Ar is an optionally-substituted aryl or an optionally-substituted heteroaryl;

provided that when the group represented by the formula (A2) is a group represented by the
formula:

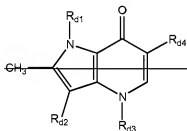


wherein R¹ is hydrogen, chloro or an optionally-substituted alkoxy and R¹ is as defined above;
and W is a bond, then Ar is not thiazolyl-substituted with one or two substituents or condensed
with dihydroimidazole;
and excluding excluding the following compounds:
(i) a compound represented by the formula:



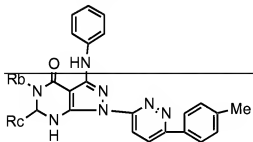
wherein Ra is a substituted carbamoyl;

(ii) a compound represented by the formula:



wherein R_{d1} and R_{d2} is each hydrocarbyl, R_{d3} and R_{d4} is each carboxy optionally substituted with hydrocarbyl;

(iii) a compound represented by the formula:



wherein Rb is hydrogen, amino or phenyl, Rc is C₁₋₄ alkyl, a substituted phenyl or an optionally substituted heteroaryl;

(iv) ethyl 4-(6-chloro-2,2,4-trimethyl-3,4-dihydro-2H-1,4-benzoxazin-8-yl)-6-propyl-2,4-dihydro-1H-pyrazolo[3,4-b]pyridine-5-carboxylate, 7-methoxy-3-(4-methoxyphenyl)-1-methyl-5-phenylquinolin-4(1H)-one, 8-methoxy-3-(4-methoxyphenyl)-1-methyl-5-phenylquinolin-4(1H)-one, 4-(8-benzyl-4-methyl-3,4-dihydro-2H-1,4-benzoxazin-6-yl)-2,4-dioxobutanoic acid, ethyl 1,7-dimethyl-4-oxo-3,5-diphenyl-1,2,3,4-tetrahydroquinazoline-6-carboxylate, 1-cyclobutyl-6,8-difluoro-7-(4-methylpiperazin-1-yl)-4-oxo-5-phenoxy-1,4-dihydroquinoline-3-carboxylic acid, 1-cyclopropyl-7-(2,6-dimethylpyridin-4-yl)-6,8-difluoro-4-oxo-5-(phenylthio)-1,4-dihydroquinoline-3-carboxylic acid, 1-ethyl-8-methoxy-5-phenylquinolin-4(1H)-one, 1-

cyclopropyl-6,8-difluoro-7-(4-methylpiperazin-1-yl)-4-oxo-5-(phenylthio)-1,4-dihydroquinoline-3-carboxylic acid, 4,6-dimethyl-8-(4-methyl-6-oxo-1,4,5,6-tetrahydropyridazin-3-yl)-2H-1,4-benzoxazin-3(4H)-one, 4,6-dimethyl-8-(6-oxo-1,4,5,6-tetrahydropyridazin-3-yl)-2H-1,4-benzoxazin-3(4H)-one, 2,2,4-trimethyl-8-(6-oxo-1,4,5,6-tetrahydropyridazin-3-yl)-2H-1,4-benzoxazin-3(4H)-one, 8-chloro-1-methyl-4-oxo-5-phenyl-1,4-dihydroquinoline-3-carboxylic acid, 8-[(4,6-dimethoxypyrimidin-2-yl)sulfinyl]-4-methyl-2-phenylphthalazin-1(2H)-one, 3-[(1,5-dimethyl-3-oxo-2-phenyl-2,3-dihydro-1H-pyrazol-4-yl)amino]-6-methyl-1,7-dihydro-4H-pyrazolo[3,4-d]pyrimidin-4-one, 6-(4-bromophenyl)-1-(4-methoxyphenyl)-5-methyl-7-oxo-6,7-dihydro-1H-pyrazolo[4,3-d]pyrimidine-3-carbonitrile, 3,6-dibenzyl-1-cyclopentyl-1,7-dihydro-4H-pyrazolo[3,4-d]pyrimidin-4-one, methyl (6-tert-butoxy-4-oxo-1,3-diphenyl-1,4-dihydro-5H-pyrazolo[3,4-d]pyrimidin-5-yl)acetate, 1,3,6-trimethyl-5-phenyl-1H-pyrrolo[2,3-d]pyrimidine-2,4(3H,7H)-dione, ethyl 4-({2-[(2,2-dimethylpropanoyl)amino]-6-methyl-4-oxo-4,7-dihydro-1H-pyrrolo[2,3-d]pyrimidin-5-yl}thio)benzoate and methyl 4-({2-[2-amino-7-benzyl-3-(isopropoxymethyl)-4-oxo-4,7-dihydro-3H-pyrrolo[2,3-d]pyrimidin-5-yl]vinyl}benzoate; and Ar is a phenyl which is substituted with

- _____ (i) one or more C₁₋₈ alkyl which may be substituted with a halogen,
- _____ (ii) one or more alkoxy,
- _____ (iii) one or more halogen,
- _____ (iv) one or more benzyloxy, or
- _____ (v) one or more hydroxy;

or a salt thereof.

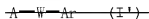
2-14. (Cancelled)

15. (Currently Amended) The compound according to claim 1, wherein the compound is 3-(2,4-dimethylphenyl)-6-dipropylamino-1,5-dimethyl-1,5-dihydro-4H-pyrazolo[3,4-d]pyrimidin-4-one, 5-(2,4-dimethylphenyl)-3-methyl-1-(1-propylbutyl)quinolin-4(1H)-one, 1-(dipropylamino)-6-mesityl-3-methyl-4H-quinolizin-4-one, 2-(dipropylamino)-5-mesityl-3,7-dimethyl-3,7-dihydro-4H-pyrrolo[2,3-d]pyrimidin-4-one;

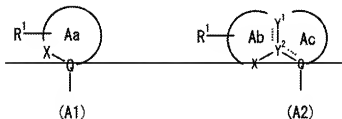
1-(2,4-dimethylphenyl)-4-(1-ethylpropoxy)-6-methyl-1,6-dihydro-7H-pyrrolo[2,3-d]pyridazin-7-one;

5-mesityl-3-methyl-1-(1-propylbutyl)cinolin-4(1H)-one, or 1-(1-ethylpropyl)-4-mesityl-2-methyl-1,2-dihydro-3H-indazol-3-one.

16. (Currently Amended) A method for treating or preventing a disease wherein a CRF receptor is implicated, which comprises administering to a subject in need thereof an effective amount of a compound or salt according to claim 1, wherein the disease being treated or prevented is selected from the group consisting of affective disorder, depression and anxiety represented by the formula:



wherein, A is a group represented by the formula (A1) or (A2):



wherein, ring Aa is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Q and X, and may be further substituted with one or more substituents in addition to R¹; ring Ab is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Y¹, Y² and X, and may be further substituted with one or more substituents in addition to R¹; ring Ac is a 5- or 6- membered ring which may have one or two further heteroatoms selected from oxygen, sulfur and nitrogen at a position other than Y¹, Y² and Q, and may be substituted with one or more substituents; R¹ is an optionally substituted alkyl, an optionally substituted cycloalkyl, an optionally substituted cycloalkenyl, a substituted amino, an optionally substituted cyclic amino, a substituted hydroxy, a substituted sulfonyl, an optionally substituted sulfinyl, or an optionally substituted sulfonyl; X is carbonyl, -O-, -S-, -SO-, or -SO₂-; Y¹, Y² and Q are independently optionally substituted carbon or nitrogen; --- is a single or double bond;

~~W is a bond, an optionally substituted methylene, an optionally substituted ethylene, an optionally substituted imino, O, S, SO, or SO₂;~~

~~Ar is an optionally substituted aryl or an optionally substituted heteroaryl;
or a salt thereof or a prodrug thereof.~~

17. (Cancelled)

18. (Currently amended) A ~~medicine~~pharmaceutical composition comprising the compound according to claim 1 or a ~~prodrug salt~~ thereof.

19. (Currently amended) The ~~medicine~~pharmaceutical composition according to claim 18 which is a corticotropin releasing factor antagonist.

20. (Currently amended) The ~~medicine~~pharmaceutical composition according to claim 18 which is an agent for treating or preventing affective disorder, depression or anxiety.

21. (Cancelled)

22. (New) The compound according to claim 1, wherein R^{1a} is

(1) an amino which is mono- or di-substituted with

(i) a C₁₋₈ alkyl which may be substituted with a methoxy, a cyclopropyl, a phenyl, a 4-methylphenyl, a phenoxy, a 2-chlorophenyl, a pyridyl, a 4-chlorophenyl, a 4-(benzyloxy)phenyl, a 3-methoxyphenyl, a 3-chlorophenyl, a 2'-cyanobiphenyl, a pyrrolyl, a naphthyl, a 2,5-dimethoxyphenyl, a quinolinyl, a 3-fluoro-5-(trifluoromethyl)phenyl, a benzoyl, an ethoxycarbonyl, or an *N,N*-dimethylcarbamoyl,

(ii) a C₂₋₈ alkenyl,

(iii) a C₁₋₁₀ acyl, or

(iv) a C₃₋₇ cycloalkyl,

(2) a piperidinyl,

(3) a pyrrolidinyl, or

(4) a morpholinyl.

23. (New) The compound according to claim 1, wherein R^{1a} is an amino which is mono- or di-substituted with a C_{1-8} alkyl.

24. (New) The compound according to claim 1, wherein R^2 is a C_{1-8} alkyl.

25. (New) The compound according to claim 1, wherein $R^{2'}$ is a C_{1-8} alkyl.

26. (New) The compound according to claim 1, wherein Ar is a phenyl which is substituted with one or more C_{1-8} alkyl.

27. (New) The compound according to claim 1, wherein R^{1a} is an amino group which is mono- or di-substituted with a C_{1-8} alkyl;

R^2 is a C_{1-8} alkyl;

$R^{2'}$ is a C_{1-8} alkyl; and

Ar is a phenyl which is substituted with one or more C_{1-8} alkyl.